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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/918,039	07/30/2001	Yong Mi Choi-Sledeski	P24450-E US1	3370
7590 12/02/2004		EXAMINER		
Synnestvedt &	Lechner LLP		TRUONG, TAN	итном NGO
2600 Aramark Tower		ART UNIT	PAPER NUMBER	
Philadelphia, PA 19107-2950			1624	
			DATE MAILED: 12/02/200	

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)			
Office Action Summary		09/918,039	CHOI-SLEDESKI ET AL.			
		Examiner	Art Unit			
		Tamthom N. Truong	1624			
	The MAILING DATE of this communication app	pears on the cover sheet with the c	correspondence address			
THE - Exte after - If the - If NO - Failu	MORTENED STATUTORY PERIOD FOR REPLIMAILING DATE OF THIS COMMUNICATION. ensions of time may be available under the provisions of 37 CFR 1.1 r SIX (6) MONTHS from the mailing date of this communication. e period for reply specified above is less than thirty (30) days, a replimate to reply within the set or extended period for reply will, by statute	136(a). In no event, however, may a reply be timely within the statutory minimum of thirty (30) days will apply and will expire SIX (6) MONTHS from e, cause the application to become ABANDONE	nely filed s will be considered timely. I the mailing date of this communication. 10 (35 U.S.C. § 133).			
earn	reply received by the Office later than three months after the mailing ned patent term adjustment. See 37 CFR 1.704(b).	g date of this communication, even if timely liled	I, may reduce any			
Status						
· -	Responsive to communication(s) filed on 20 A	ugust 2004.				
2a)□	•—	s action is non-final.				
3)□	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.					
Dispositi	ion of Claims					
5) 6) 7)	Claim(s) 1,2,6,8-12,14,22 and 24-41 is/are per 4a) Of the above claim(s) is/are withdraw Claim(s) is/are allowed. Claim(s) is/are rejected. Claim(s) is/are objected to. Claim(s) 1, 2, 6, 8-12, 14, 22, and 24-41 are su	wn from consideration.	n requirement.			
Applicati	ion Papers					
9) The specification is objected to by the Examiner.						
10)	10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.					
	Applicant may not request that any objection to the	drawing(s) be held in abeyance. See	37 CFR 1.85(a).			
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d). 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority u	under 35 U.S.C. § 119					
a)[Acknowledgment is made of a claim for foreign All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the priority documents application from the International Bureau See the attached detailed Office action for a list of	s have been received. s have been received in Application rity documents have been received u (PCT Rule 17.2(a)).	on No ed in this National Stage			
Attachment	t(s)					
	e of References Cited (PTO-892)	4) Interview Summary ((PTO-413)			
3) 🔲 Inform	e of Draftsperson's Patent Drawing Review (PTO-948) mation Disclosure Statement(s) (PTO-1449 or PTO/SB/08) r No(s)/Mail Date	Paper No(s)/Mail Dat				

DETAILED ACTION

Applicant's amendment of 8-20-04 has been considered. Due to the extensive number of rings or ring systems present, the following restriction is required. The previous 112, and 102 rejections are set aside right now, and will be addressed once an election is made.

Claims 3-5, 7, 13, 15-21, and 23 are cancelled.

Claims 1, 2, 6, 8-12, 14, 22, and 24-41 are pending.

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

1. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I, wherein:

Ar¹ is pyrrolo[2,3-c]pyridinyl;

R₂ is SO₂-phenyl, or SO₂-naphthyl;

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

2. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[2,3-c]pyridinyl;

 R_2 is SO_2 -(5-membered heteroaryl or heterocyclyl);

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Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

3. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I, wherein:

Ar¹ is pyrrolo[2,3-c]pyridinyl;

 R_2 is SO_2 -(6-membered heteroaryl)

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

4. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[2,3-c]pyridinyl;

R₂ is SO₂-quinolinyl;

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

5. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[2,3-c]pyridinyl;

R₂ is SO₂-benzopyranyl;

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Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

6. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I, wherein:

Ar¹ is pyrrolo[2,3-b]pyridinyl;

R₂ is SO₂-phenyl, or SO₂-naphthyl;

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

7. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[2,3-b]pyridinyl;

R₂ is SO₂-(5-membered heteroaryl or heterocyclyl);

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

8. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I, wherein:

Ar¹ is pyrrolo[2,3-b]pyridinyl;

R₂ is SO₂-(6-membered heteroaryl)

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Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

9. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[2,3-b]pyridinyl;

R₂ is SO₂-quinolinyl;

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

10. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[2,3-b]pyridinyl;

 R_2 is SO_2 -benzopyranyl;

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

11. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I, wherein:

Ar¹ is pyrrolo[3,2-c]pyridinyl;

R₂ is SO₂-phenyl, or SO₂-naphthyl;

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Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

12. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[3,2-c]pyridinyl;

R₂ is SO₂-(5-membered heteroaryl or heterocyclyl);

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

13. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I, wherein:

Ar¹ is pyrrolo[3,2-c]pyridinyl;

R₂ is SO₂-(6-membered heteroaryl)

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

14. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[3,2-c]pyridinyl;

R₂ is SO₂-quinolinyl;

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Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

15. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to compounds of formula I wherein:

Ar¹ is pyrrolo[3,2-c]pyridinyl;

R₂ is SO₂-benzopyranyl;

Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds;

classified in classes 514, 546, various subclasses depending on substituents.

- 16. Claims 1, 2, 6, 8-12, 14, 22, and 24-34, drawn to the remaining compounds of formula I wherein the combination of Ar¹ and R₂ is not in the above groups.

 Pharmaceutical composition thereof, and method of treating thrombus formation (inhibiting Factor Xa) using said compounds; classified in classes 514, 546, various subclasses depending on substituents. Further restriction will be required if this group is elected.
- 17. Claims 35-41, drawn to the combination therapy and/or composition comprising additional agent(s); classified in class 514, various subclasses depending on the additional agents. Further restriction will be required if this group is elected.

The inventions are distinct, each from the other because of the following reasons:

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- a. Inventions groups 1-17 are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the combinations of rings represented by Ar¹, R₂ and the pyrrolidinone ring define the different inventions.
- b. Athough all groups share the ring of *pyrrolidinone*, said ring alone does not sufficiently define the invention, and does not contribute to the art. Therefore, it is the combination of the *pyrrolidinone* with Ar¹ and R₂ that gives each group a distinct physical, chemical and/or biological properties, and thus sets apart the compounds of one group from those of the others. Thus, a reference that anticipated, or rendered obvious one group would not do so to the others, and so, a separate search is required for each group.

Because these inventions are distinct for the reasons given above and the search required for Group 1 is not required for Group 2-17, and the search for all 17 distinct invention would impose a serious burden upon the examiner in charge of this invention, restriction for examination purposes as indicated is proper.

Due to the complexity of the grouping, the restriction is presented in writing.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the

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application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Tamthom N. Truong whose telephone number is 571-272-0676. The examiner can normally be reached on M-F (10:00-6:30).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, James O Wilson can be reached on 571-272-0661. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Tamthom N. Truong

Examiner

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11-15-04

JAMES O. WILSON

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